

# Cambrian–Ordovician conodonts from slump deposits of the Argentine Precordillera: new insights into its passive margin development

GUSTAVO G. VOLDMAN\*<sup>†‡</sup>, JUAN L. ALONSO<sup>§</sup>, LUIS P. FERNÁNDEZ<sup>§</sup>,  
ALDO L. BANCHIG<sup>¶</sup>, GUILLERMO L. ALBANESI\*<sup>‡</sup>, GLADYS ORTEGA\*  
& RAÚL CARDÓ<sup>||</sup>

\*CONICET, Museo de Paleontología, CIGEA, Facultad de Ciencias Exactas, Físicas y Naturales, Universidad Nacional de Córdoba, X5016GCB Córdoba, Argentina

<sup>‡</sup>CICTERRA (CONICET-UNC), Córdoba, Argentina

<sup>§</sup>Departamento de Geología, Universidad de Oviedo, 33005 Oviedo, Spain

<sup>¶</sup>Facultad de Ciencias Exactas y Naturales, Universidad Nacional de San Juan, 5400 San Juan, Argentina

<sup>||</sup>Servicio Geológico Minero Argentino (SEGEMAR), 5400 San Juan, Argentina

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**Abstract** – The Los Sombreros Formation represents the western continental margin slope deposits of the Argentine Precordillera, a sub-terrane accreted to Gondwana as part of the Cuyania Terrane in early Palaeozoic times. The age of these gravity-driven deposits is controversial and, therefore, a precise biostratigraphic scheme is essential to reveal the evolution of the continental margin. New conodont samplings along with sedimentological and structural analysis carried out in the Los Sombreros Formation in the La Invernada Range provide clues to its depositional framework. The sedimentary succession is made up of dominantly calciturbidites, carbonate breccias and conglomerates, along with mudstones that represent the pelagic/hemipelagic background sedimentation. It displays hectometric to outcrop-scale slump folds with variable hinge-line orientations and pinch-and-swell structures, evidencing soft-sediment deformation, consistent with a slope to base-of-slope setting. Three limestone samples from this succession include conodonts referable to the pandemic *Hirsutodontus simplex* Subzone of the *Cordylodus intermedius* Zone (upper Furongian, Cambrian) and from the *Macerodus diana* Zone (upper Tremadocian, Ordovician), implying that a slope connected the shallow-water shelf with a deep-water (oceanic) basin at least since late Cambrian times. The conodont faunas show affinities to coeval assemblages from outer shelf and slope environments around Laurentia yet they are not conclusive to postulate a geographic origin for the Precordillera. The thermal alteration of the conodonts is consistent with sedimentary burial and nappe stacking in this sector of the Precordillera.

Keywords: conodont, slope facies, slump, Los Sombreros Formation, Argentine Precordillera, Cambrian, Ordovician.

## 1. Introduction

In the external zone of the Andean orogen, the Argentine Precordillera records a series of tectono-stratigraphic events related to the building of Gondwana during early to middle Palaeozoic times (e.g. Cawood, 2005; Rapela *et al.* 2016). Eastern, Central and Western domains have classically been distinguished after their structural and stratigraphic features (Ortiz & Zambrano, 1981; Baldis *et al.* 1982). The Eastern and Central Precordillera involve a large passive-margin carbonate platform, Cambro-Ordovician in age, which is overlain by Middle Ordovician siliciclastic foreland-basin deposits that reach up to the Devonian Punta Negra Formation. Conversely, the Western Precordillera exhibits a deep-water succession, with ocean-floor sedimentary rocks containing pillow lavas and mafic–ultramafic bodies in the westernmost sections.

The transition between the Central and Western Precordillera is represented by the disorganized deposits of the Los Sombreros Formation (Cuerda, Cingolani & Varela, 1983; Banchig, Keller & Milana, 1990) and the Corralito Formation (Furque & Caballé, 1988; Furque *et al.* 1990), recording the slope of a continental margin. However, the complex structure and the scarcity of fossils challenge against a precise depositional scheme, whilst Ordovician and Devonian ages have been proposed for the Los Sombreros Formation (e.g. Benedetto & Vaccari, 1992; Voldman, Albanesi & Ramos, 2009; Peralta, 2013). For instance, Peralta (2013) considered that all the disorganized deposits slid in Devonian times, after sedimentation of the Punta Negra Formation. Thus, constraining the spatio-temporal framework of the mélanges is essential for understanding the geotectonic evolution of the passive continental margin of the early Palaeozoic Precordillera.

In the present study, new conodont findings along with structural, stratigraphic and sedimentological data

<sup>†</sup>Author for correspondence: [gvoldman@unc.edu.ar](mailto:gvoldman@unc.edu.ar)

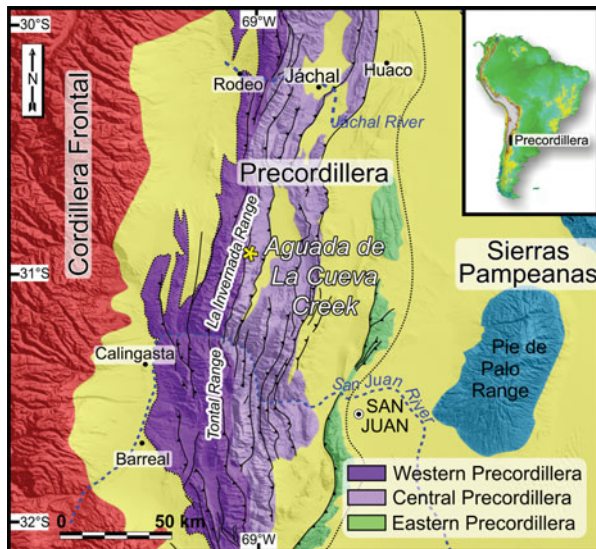


Figure 1. (Colour online) Location map of the Argentine Precordillera fold-and-thrust belt showing the studied locality and the western, central and eastern morphostructural domains. Base image derived from Shuttle Radar Topography Mission (SRTM).

72 for the Los Sombreros Formation in the La Invernada Range are presented. The record of the *Hirsutodontus simplex* Subzone of the *Cordylodus intermedius* Zone (upper Furongian, Cambrian) and the *Macerodus dianae* Zone (upper Tremadocian, Ordovician) in gravity-flow deposits with syndepositional deformation features suggests the existence of a slope in late Cambrian – Early Ordovician times. Moreover, the defined high-resolution conodont biostratigraphy improves the intrabasinal correlation with the Precordilleran carbonate platform as well as with other regions of the world.

## 84 2. Geological setting

85 The Argentine Precordillera of La Rioja, San Juan and Mendoza provinces is a *c.* 80 km wide foreland fold-and-thrust belt that involves Palaeozoic to Cenozoic rocks, which extends between 29° and 33° S in the Andean foothills above the shallow subduction segment of the Nazca plate (Heim, 1952; Allmendinger *et al.* 1990; Gosen, 1992; Ramos, Cristallini & Pérez, 2002) (Fig. 1). The Precordilleran Cambrian–Ordovician carbonate platform sequence is unique to South America and makes up part of a larger region of the Andean foothills of western Argentina that is referred to as the Cuyania composite terrane (Ramos, 1995).

97 At the continental slope and rise, to the west of the Precordilleran carbonate platform, the Los Sombreros mélangé is a mudstone-dominated deep-water disorganized unit, containing a diverse array of rocks derived from the platform, the slope and the basement. These rocks include arkosic sandstones, megabreccias, conglomerates with well-rounded basement-derived metamorphic and igneous pebbles to boulders, thin-bedded fine-grained limestones, carbonate breccias and blocks up to several hectometres in size of lower Cambrian to

107 Lower Ordovician limestones displaying platform facies not represented elsewhere (e.g. Bordonaro, 2003).

108 The Los Sombreros Formation displays ubiquitous extensional structures that result in block-in-matrix fabric in some places as a consequence of submarine sliding (Alonso *et al.* 2008). The main outcrop belt of the mélangé extends along the eastern flank of the Tontal Range (Fig. 1), where the formation is more than 1000 m thick in the Seca Creek type section (Cuerda, Cingolani & Varela, 1983; Cuerda *et al.* 1986). The outcrops continue patchily to the north up to the Jáchal River area (Benedetto & Vaccari, 1992), whereas its southern prolongation is represented by the Estancia San Isidro Formation in Mendoza, which exhibits giant Cambrian limestone blocks enclosed in a green shaly matrix of Darriwilian age (Keller, 1999; Heredia & Beresi, 2004; Ortega *et al.* 2007). Thus, the slope facies of the Precordillera extends over 300 km with N–S orientation (Fig. 1).

## 126 3. Previous biostratigraphic studies of the Los Sombreros Formation

128 The Los Sombreros Formation was originally referred to the Lower Ordovician Series based on graptolite records from its lower third (Cuerda *et al.* 1985). Later trilobite findings revealed the presence of lower and middle Cambrian rocks (Bordonaro & Baldis, 1987; Bordonaro & Banchig, 1990), interpreted as out-platform resedimented blocks in a Lower – Middle Ordovician succession with autochthonous conodonts and graptolites (Benedetto & Vaccari, 1992; Bordonaro, 2003). Lehnert (1994) described the first conodont assemblage of the *Cordylodus proavus* Zone (upper Furongian) in the Precordillera, from outcrops of the Los Sombreros Formation in the Tontal Range. The conodont association includes *Cordylodus primitivus* Bagnoli, Barnes & Stevens, *Cordylodus proavus* Müller and *Eoconodontus notchpeakensis* Miller. These conodont elements derive from a calcisiltite lens that overlies shales with graptolite specimens that demonstrate its allochthonous character.

147 Albanesi, Ortega & Hünicken (1995) proposed that the conformable stratigraphic contact between the Los Sombreros Formation and the overlying Yerba Loca Formation at Ancaucha Creek, 10 km northwest of Jáchal city, is early Darriwilian in age. They determined that the top of the Los Sombreros Formation at this locality is lower Darriwilian at the most, by considering the presence of the index conodont *Baltoniodus clavatus* Stouge & Bagnoli from the basal beds of the Yerba Loca Formation and the absence of the key species *Paroistodus horridus* (Barnes & Poplawski), which appears *c.* 40 m above the Yerba Loca base.

159 Further fossil findings include upper Cambrian, Tremadocian, Floian and Darriwilian conodont faunas (Voldman, Albanesi & Ramos, 2009; Voldman *et al.* 2014) as well as graptolites referable to the uppermost Tremadocian, Floian and Sandbian stages (e.g. A. L. Banchig, unpub. Ph.D. thesis, Univ. Nacional San Juan,



Figure 2. (Colour online) Geological map of the study area with fossiliferous sampling points.

1995; Banchig & Moya, 2002; Ortega *et al.* 2014). Additionally, Astini, Thomas & Yochelson (2004) identified the enigmatic fossil *Salterella maccullochi* (Murchison) in the Ancaucha Olistolith of the Los Sombreros Formation, composed of inner-shelf deposits that they interpreted as lower Cambrian synrift strata.

4. Study area and methods

The La Invernada Range constitutes a critical region to investigate the architecture and kinematic development of the disorganized deposits of the Los Sombreros Formation. This range lies to the north of the Tontal Range, extending for *c.* 60 km with a N–S trend (Fig. 1). It displays E-verging thrusts and related folds, involving a lower Palaeozoic succession that also includes the Siluro-Devonian mélangé deposits of the Corralito Formation, which crop out exclusively in this range. In order to determine the age of the Los Sombreros Formation at the Aguada de La Cueva Creek section (Fig. 2), nine limestone samples (14 kg in total) were collected and processed following the standard techniques to recover conodont elements (Stone, 1987). Three samples were productive, yielding 80 specimens that are housed under the repository codes CORD-MP 50734 to 50814 in the Museo de Paleontología, Facultad de Ciencias Exactas, Físicas y Naturales, Universidad Nacional de Córdoba, Argentina. Field work also included structural mapping and stratigraphic and sedimentological studies, during which kinematic data were acquired and rocks were described in terms of their bedding and lithology and sampled for thin-section preparation.

5. Results

5.a. Stratigraphy and structure

In the Aguada de La Cueva Creek area, the Los Sombreros Formation consists of a *c.* 500–600 m thick succession of dominantly limestones, carbonate breccias and conglomerates and subordinate shales (Fig. 2). The limestones are thin bedded, varying from peloidal/calcsphere mudstones and wackestones to graded and laminated calcilithites (calciturbidites) (Fig. 3a, b), which may pass upwards into a shale divi-

sion and gradationally overlie a breccia/conglomerate division. Breccias/conglomerates contain mainly clasts of mudstones, peloidal grainstones and recrystallized limestones, with minor amounts of chert and quartz clasts. They constitute beds that are massive or graded, and that may be gradationally overlain by a calciturbidite; some exceptional examples of stratified (laminated) beds also occur (Fig. 3c). The whole succession lacks *in situ* shallow-water faunas, with the exception of scarce lingulids, and exhibits slight bioturbation restricted to the peloidal/calcsphere limestones. Remarkably, it displays hectometric slump folds, extensional faults and lateral transitions between intact and faulted/brecciated beds (Fig. 3a, d).

As a whole, most of these facies are indicative of sedimentation from gravity flows, from cohesive debris flows to low-density turbidity currents (see Lowe, 1982; McIlreath & James, 1984; Mutti *et al.* 1999, amongst others). Fine-grained, peloidal/calcsphere mudstones probably represent the pelagic/hemipelagic background sedimentation. This interpretation is compatible with the synsedimentary deformation features (see also Section 5.a.1. below), all indicating a slope to base-of-slope setting. The relatively deep-water environment is also suggested by the absence of *in situ* shallow-water fauna and the scarcity of bioturbation.

The slumped succession at the Aguada de La Cueva Creek section is unconformably overlain by grey shales with poorly preserved graptolites, including *Archiclimacograptus* sp., *Dicellograptus* sp., *Dicranograptus* sp., *Nemagraptus* sp. and *Reteograptus speciosus* Harris, which suggest a late Darriwilian to Sandbian age for the upper part of the Los Sombreros Formation in this area (Sample AMGRAP, Figs 2, 4). These graptolitic shales are in turn unconformably overlain by sandstone–mudstone alternations with scarce calcarenites and interbedded conglomerates and calcareous breccias of the Sierra de La Invernada Formation (Furque *et al.* 1990), which are Middle–Late Ordovician in age (Ortega *et al.* 2008).

The Los Sombreros Formation is carried on a W-dipping thrust surface onto the Siluro-Devonian Corralito mélangé (Fig. 2), which comprises greenish grey shales and coquinas with extensional features, such as boudins, as well as limestone blocks of the San Juan Formation. The thrust surface displays *C'*-type shear bands and striations, recording an eastward movement of the hanging wall. It is probably Andean in age because it is located at the toe of the Invernada Range and therefore should have been responsible for the uplift of this modern geomorphic feature. However, this thrust could be the result of rejuvenation of an older Chanic or Gondwanic thrust during Andean times, which commonly occurs in the Argentine Precordillera (Ramos, Vujovich & Dallmeyer, 1996; Alonso *et al.* 2005).

5.a.1. Synsedimentary deformation

Regarding the internal structure of the Los Sombreros Formation, the most conspicuous features in the study

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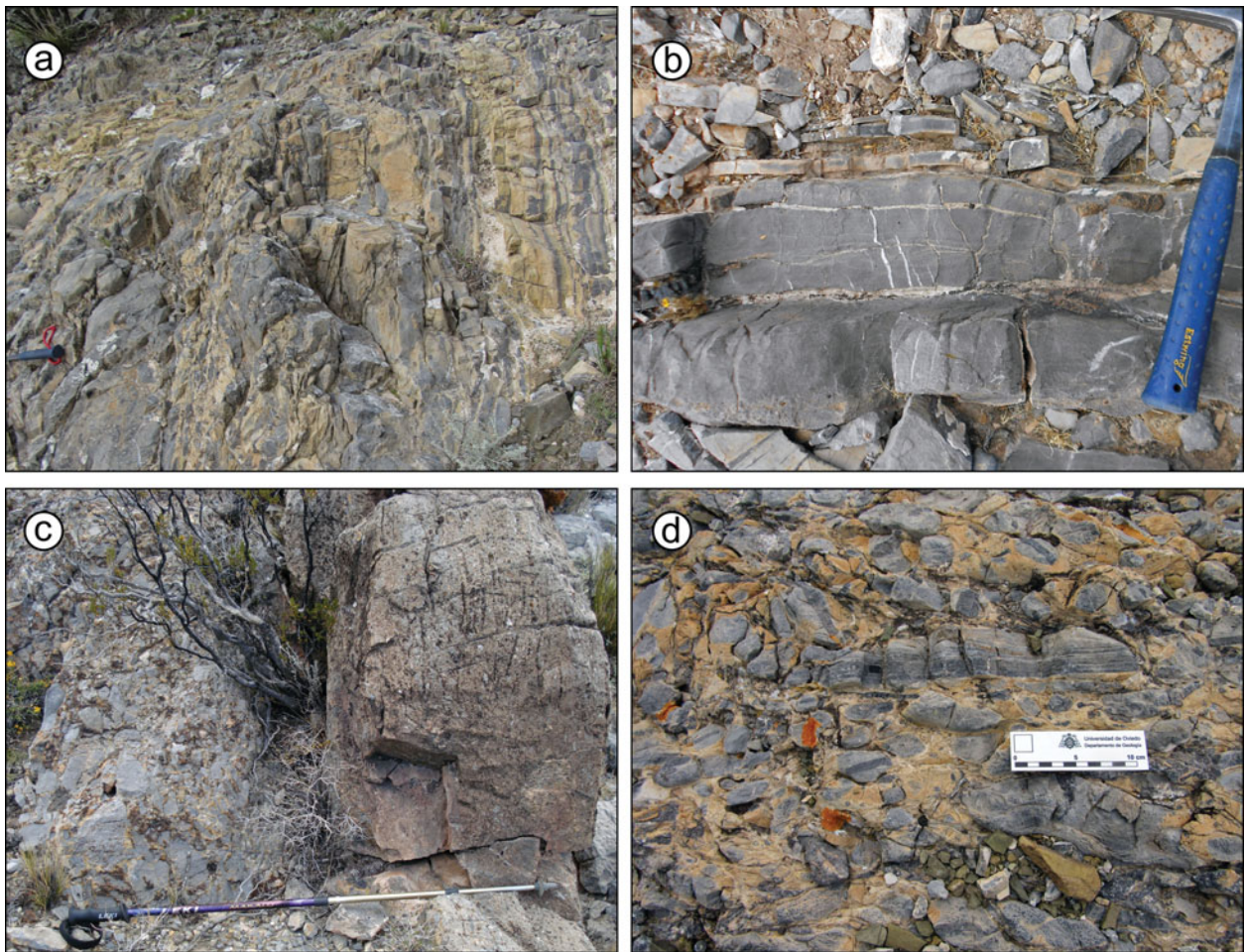


Figure 3. (Colour online) Field photographs of selected deposits from the Los Sombreros Formation at the Agua de la Cueva Creek. (a) Detail of an interval made of pebble-to-cobble breccias (left of photograph) and thin-bedded limestones (laminated mudstones and graded and laminated calciturbidites). Notice the local disruption of bedding, which laterally passes into undisturbed bedding. Stratigraphic top towards the left. (b) Interval formed of calciturbidites and variably laminated, slightly bioturbated mudstones with peloids and calcispheres. The lowermost bed is a graded calciturbidite evolving from a basal fine-pebble-bearing granulestone to a sand-grade division, whose upper part is faintly laminated and displays smooth undulations. Younging direction is to the top of the photograph. Hammer handle for scale *c.* 19 cm long. (c) Poorly sorted boulder-to-pebble carbonate breccia/conglomerate, displaying two divisions, a lower graded division and an upper laminated division. Walking pole is *c.* 120 cm long. Stratigraphic top towards the right. (d) Close-up of an interval made of thin-bedded lime mudstones, commonly laminated, with a variable degree of folding and bed disruption and brecciation. Bedding in the area is parallel to the bed above the scale, which is almost intact. The yellowish matrix surrounding the clasts is mainly dolomitic. Younging direction is to the top of the photograph.

264 area are hectometric folds that can be interpreted as  
 265 slump structures (Fig. 2). These folds do not involve  
 266 the overlying Sierra de La Invernada Formation. As  
 267 well, most of the smaller, outcrop-scale structures are  
 268 slump folds with variable hinge-line orientations and  
 269 pinch-and-swell structures, which record soft-sediment  
 270 deformation (Fig. 5). Tension fractures perpendicular  
 271 to bedding and normal faults also occur. In the ex-  
 272 ample of Figure 5a, b, tension fractures are restricted  
 273 to the yellowish beds, indicating that these beds under-  
 274 went brittle behaviour, while other beds, with pinch-  
 275 and-swell boudinage, were stretched by more ductile  
 276 deformation, probably because the yellowish beds were  
 277 more lithified. So, pinch-and-swell boudins and tension  
 278 and shear fractures are more or less coeval and all of  
 279 them imply bed-parallel extension. Although the devel-  
 280 opment of the slump fold in Figure 5a, b is prior to ex-  
 281 tensional deformation (both fold limbs are truncated by

a normal fault), all the above mentioned structures sup-  
 port the interpretation that gravitational collapse and  
 sliding was the cause of the deformation. In Figure 5d,  
 the cut-off lines of the tension fractures (blue) are par-  
 allel to the boudin necks lineation (red), recording the  
 same extension direction.

### 5.b. Conodont fauna and biostratigraphy

Three carbonate samples taken at the Aguada de  
 La Cueva Creek section yielded conodonts, which  
 constrain the age of the slump deposits of the Los  
 Sombreros Formation with a high-resolution biostrati-  
 graphy and improve the conodont biozonation scheme  
 of the Precordillera. The taxonomy of the identified spe-  
 cies is well known, following descriptions of previous  
 authors; therefore, only a brief discussion is presented  
 herein.

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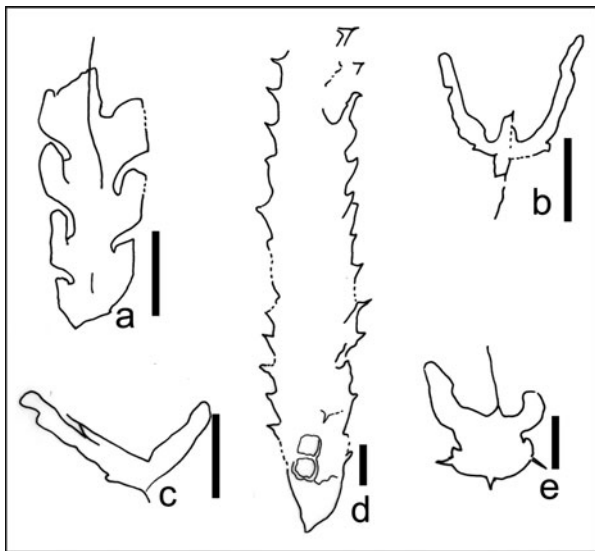


Figure 4. Late Darriwilian to Sanbian graptolites from the upper levels of the Los Sombreros Formation (Sample AMGRAP, see location in fig. 2). (a) *Archiclimacograptus* sp., CORD-PZ 25705; (b) *Dicellograptus* sp., CORD-PZ 25706; (c) *Reteograptus speciosus* Harris, CORD-PZ 25707; (d) *Nemagraptus* sp., CORD-PZ 25708; (e) *Dicranograptus?* sp., CORD-PZ 25709. Scale bar: 1 mm.

A lime mudstone affected by syndimentary extensional faults (sample AM6) yielded *Cordylodus caboti* Bagnoli, Barnes & Stevens, *C. cf. tortus* Barnes, *C. intermedius* Furnish, *C. proavus* Müller, *C. cf. andresi* Viira & Sergeyeva, *Drepanoistodus* sp., *Teridontus nakamurai* (Nogami), *Variabiloconus datsonensis* (Druce & Jones), *Westergaardodina* sp. and the index species *Hirsutodontus simplex* (Druce & Jones) (Fig. 6). The latter species is chronostratigraphically restricted to the *Cordylodus intermedius* Zone, Stage 10 of the Furongian. *H. simplex* is characterized by a simple cone with circular cross-section and a series of spines scattered mainly on the anterior and lateral sides of the base and cusp (Fig. 6n).

The different species of *Cordylodus* were distinguished by considering the general shape of the elements, the pattern of denticulation (discrete, confluent) and the basal cavity configuration (number of apices, depth, position and shape of its anterior border). As described by Bagnoli, Barnes & Stevens (1987), the basal cavity of *C. caboti* (Fig. 6g–i) is not as deep as in *C. proavus* (Fig. 6a–e), but also extends above the posterior process. Its basal cavity displays a slightly concave to straight anterior margin that

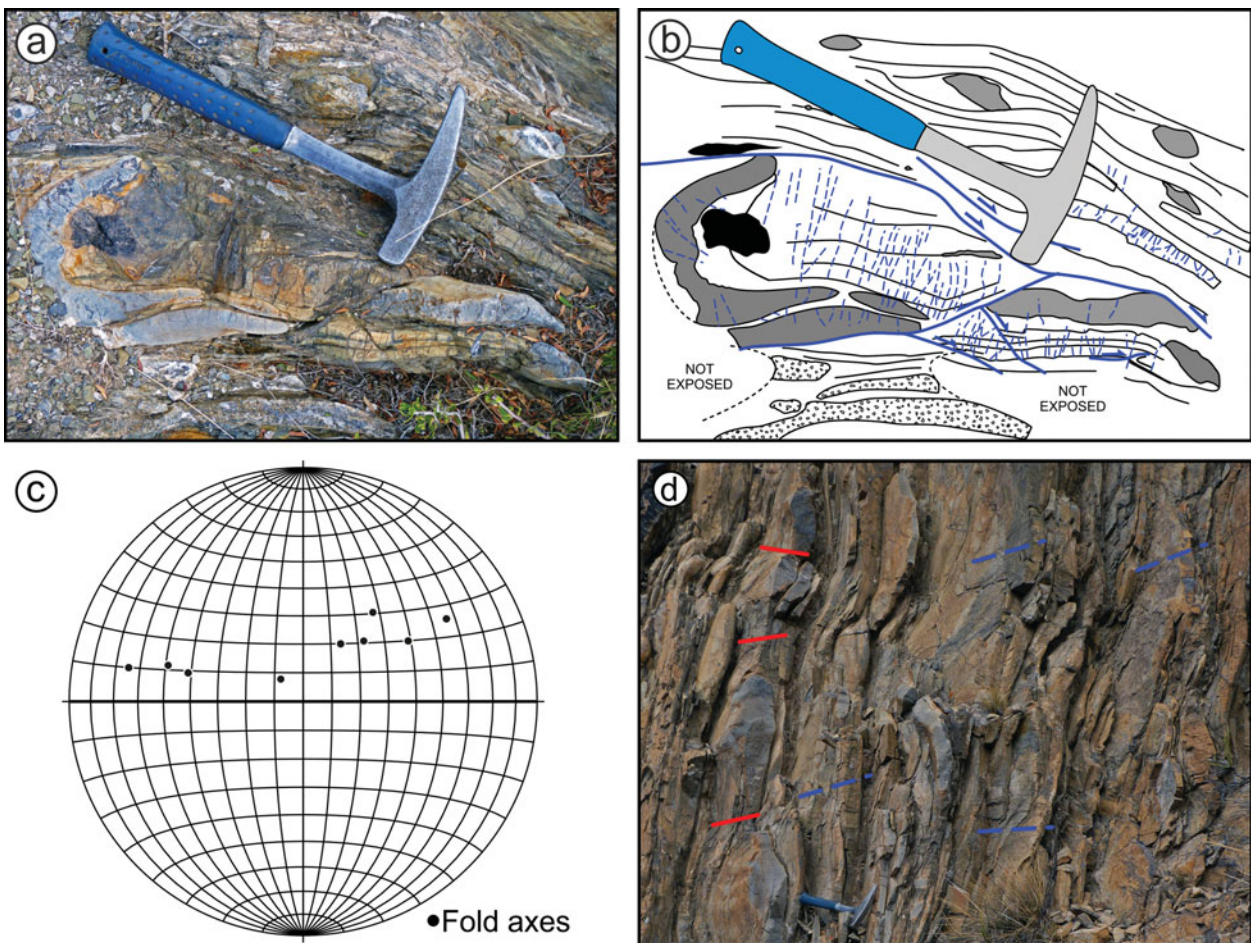


Figure 5. (Colour online) (a, b) Photograph and outcrop sketch showing a slump fold and pinch-and-swell structures in the carbonate succession of the Los Sombreros Formation. Tensional and shear fractures are depicted. See text for explanation, and location in Figure 2. (c) Stereonet of fold axes in the study area. (d) Boudinaged limestone bed (on the left) and closely spaced microfractures (on the beds located to the right). Hammer for scale: 33 cm.

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Figure 6. (Colour online) Conodont elements from the *Hirsutodontus simplex* Subzone of the *Cordylodus intermedius* Zone (upper Furongian, Cambrian), sample AM6: (a–f) *Cordylodus proavus* Müller, (a) S element, lateral view, CORD-MP 50734; (b) S element, lateral view, CORD-MP 50735; (c) S element, lateral view, CORD-MP 50736; (d) P element, lateral view, CORD-MP 50737; (e) S element, lateral view, CORD-MP 50738; (f) S element, lateral view, CORD-MP 50739. (g–i) *Cordylodus caboti* Bagnoli, Barnes & Stevens, (g) S element, lateral view, CORD-MP 50740; (h) S element, lateral view, CORD-MP 50741; (i) S element, lateral view, CORD-MP 50742. (j) *Cordylodus cf. tortus* Barnes, S element, lateral view, CORD-MP 50743. (k–m, o, p) *Cordylodus intermedius* Furnish, (k) S element, lateral view, CORD-MP 50744; (l) S element, lateral view, CORD-MP 50745; (m) S element, lateral view, CORD-MP 50746; (o) M element, lateral view, CORD-MP 50747; (p) S element, lateral view, CORD-MP 50748. (n) *Hirsutodontus simplex* (Druce & Jones), CORD-MP 50749, (n1) anterolateral view, (n2) posterior view. (q) *Westergaardodina* sp., lateral view,

322 curves near the tip, with its apex centrally located,  
 323 which differentiates it from *C. intermedius* (Nicoll,  
 324 1991; Pyle & Barnes, 2002; Zeballo & Albanesi,  
 325 2009). Miller *et al.* (2003) regarded *C. caboti* as a  
 326 junior synonym of *Cordylodus 'drucei'* Miller; since  
 327 we cannot follow his criteria in our collection, we  
 328 maintain the species *C. caboti* as valid.

329 The *Cordylodus intermedius* Zone is divided into  
 330 a lower *Hirsutodontus simplex* Subzone and an upper  
 331 *Clavohamulus hintzei* Subzone (Miller *et al.* 2003). The  
 332 lower subzone begins with the First Appearance Datum  
 333 (FAD) of *H. simplex* whereas its top is defined by the  
 334 FAD of *C. hintzei*. The presence of advanced species  
 335 of *Cordylodus* in sample AM6, such as *C. caboti* and  
 336 *C. intermedius*, which are more frequent in the upper  
 337 subzone, in the absence of *C. hintzei*, suggests an upper  
 338 *H. simplex* Subzone for this stratigraphic level (e.g.  
 339 Ross *et al.* 1997).

340 *Cordylodus proavus* extends from the upper Cam-  
 341 brian up to the *Iapetognathus* Zone, indicative of the  
 342 base of the Ordovician, whereas *C. intermedius* and  
 343 *C. caboti* reach up to the *Rossodus manitouensis* Zone  
 344 (Pyle & Barnes, 2002). Two elements recovered com-  
 345 pare well to the figured forms of *Cordylodus cf. andresi*  
 346 *sensu* Zeballo & Albanesi (2009) (Fig. 6x), which ex-  
 347 hibit a narrower cavity compared to the nominal species  
 348 (re-illustrated by Miller *et al.* 2015). Additionally, *C.*  
 349 *andresi* occurs in older rocks as it is restricted to the  
 350 *Hirsutodontus hirsutus* and *Fryxellodontus inornatus*  
 351 zones of the *C. proavus* Zone. *Cordylodus cf. tortus* is  
 352 strongly asymmetric, with a denticle flexed compared  
 353 to the cusp plane and a basal cavity relatively shallow  
 354 with its apex slightly displaced to the posterior region  
 355 (Zeballo & Albanesi, 2009).

356 The taxonomy of simple cones is complex, particu-  
 357 larly owing to their high variability and subtle diag-  
 358 nostic features. *Variabiliconus* shares with *Teridontus*  
 359 a similar apparatus plan, an abrupt junction between the  
 360 hyaline base and albid cusp, and the surface microstri-  
 361 ation. In particular, *Variabiliconus (Oneotodus) dat-*  
 362 *sonensis* closely resembles *Teridontus nakamurai* but is  
 363 distinguished by the circular basal outline, the upturn-  
 364 ing of the oral termination of the cusp and the presence  
 365 of shallow furrows associated with carinas (Druce &  
 366 Jones, 1971; Zeballo & Albanesi, 2009). Miller (1980)  
 367 reassigned *Oneotodus datsonensis* Druce & Jones to  
 368 *T. nakamurai* and to *Semiacontiodus nogamii* Miller.  
 369 Nicoll (1994) rejected the latter suggestion as well as  
 370 the emended diagnosis given by Ji & Barnes (1994a),  
 371 as they are not applicable to the type species of the  
 372 genus, which lacks lateral grooves, costae and keels.  
 373 Tolmacheva & Abaimova (2009) suggested introducing

a new genus for hybrid *Teridontus*-like species after fur-  
 374 ther investigations. The limited number of specimens  
 375 recovered does not allow for adequately defining the  
 376 latitude of morphologic variability of these primitive  
 377 forms. Therefore, we cautiously assign a simple sub-  
 378 symmetrical coniform element with long cusp to *T. na-*  
 379 *kamurai* (Fig. 6u) and those with short, erect and keeled  
 380 cusps with weak grooves to *V. datsonensis*, which is re-  
 381 stricted to the Furongian (Fig. 6w). Younger species  
 382 of *Variabiliconus* are easier to distinguish as they ex-  
 383 hibit an increase of ornamentation and development  
 384 of the cusp (Löfgren, Repetski & Ethington 1998; Ze-  
 385 ballo & Albanesi, 2013a). The *Teridontus nakamurai*  
 386 specimen recovered from the Los Sombremos Forma-  
 387 tion lacks the typical microstriation of the genus, as  
 388 previously observed by Lehnert (1994). The genus *Or-*  
 389 *minskia* also resembles *Teridontus* and lacks microstri-  
 390 ation, yet it has a hyaline cusp (Landing, Westrop &  
 391 Keppie, 2007).

392 Two samples obtained from thin-bedded fine-grained  
 393 calciturbidites interbedded with marlstones and brec-  
 394 cias located a few metres stratigraphically above AM6  
 395 were also productive. Sample AM8A yielded one  
 396 specimen of *Macerodus diana*e Fåhraeus & Nowlan  
 397 (Fig. 6s), a distinctive form of the standard North  
 398 American Midcontinent Realm zonation (Ross *et al.*  
 399 1997), whose biostratigraphic range is restricted to  
 400 a narrow interval of the upper Tremadocian. The  
 401 *M. diana*e Zone correlates with the lower part of  
 402 the upper subzone of the *Paltodus deltififer* Subzone  
 403 of the Baltoscandian scheme (Webby *et al.* 2004).  
 404 Sample AM8B contains eight elements including *Dre-*  
 405 *panodus arcuatus* Pander, *Paltodus aff. inaequalis*  
 406 (Pander) (Fig. 6y), *Rossodus cf. manitouensis* Repet-  
 407 ski & Ethington (Fig. 6r), *Scolopodus cf. subrex* Ji  
 408 & Barnes (Fig. 6t) and a cluster of the paracon-  
 409 odont *Phakelodus tenuis* (Müller). The *S. subrex* Zone  
 410 partly correlates with the *Macerodus diana*e Zone  
 411 (Pyle & Barnes, 2002), whereas *Rossodus cf. man-*  
 412 *itouensis* points to a slightly older Ordovician age;  
 413 yet the available material is not sufficient to verify its  
 414 taxonomy.  
 415

### 5.c. Conodont palaeoecology and palaeobiogeographic considerations

416 The recognition of the Furongian *Hirsutodontus*  
 417 *simplex* Subzone of the *Cordylodus intermedius* Zone  
 418 and the Tremadocian *Macerodus diana*e Zone in  
 419 the Los Sombremos Formation allows the biostrati-  
 420 graphic correlation between the slope facies and the  
 421 carbonate-platform domain to be improved, as both  
 422  
 423

CORD-MP 50750. (u) *Teridontus nakamurai* (Nogami), S element, lateral view, CORD-MP 50751. (v) *Drepanoistodus* sp., S element, lateral view, CORD-MP 50752. (w) *Variabiliconus datsonensis* (Druce & Jones), CORD-MP 50753, (w1) lateral view, (w2) basal cavity view. (x) *Cordylodus cf. andresi* Viira & Sergeyeva, S element, lateral view, CORD-MP 50754. (r–t, y) Conodont elements from the *Macerodus diana*e Zone (upper Tremadocian, Ordovician): (r) *Rossodus cf. manitouensis* Repetski & Ethington, S element, lateral view, sample AM8B, CORD-MP 50755; (s) *Macerodus diana*e Fåhraeus & Nowlan, lateral view, sample AM8A, CORD-MP 50756; (t) *Scolopodus cf. subrex* Ji & Barnes, lateral view, sample AM8B, CORD-MP 50757; (y) *Paltodus aff. inaequalis* (Pander), lateral view, sample AM8B, CORD-MP 50758. Scale bar: 0.1 mm.

424 zones occur within the La Silla Formation (Lehnert,  
425 Miller & Repetski, 1997). In particular, Albanesi,  
426 Cañas & Mango (in press) described thoroughly the  
427 conodont association of the *M. diana* Zone for the  
428 shallow-water carbonates of the La Silla Formation, at  
429 the Cerro Viejo de San Roque section. *Hirsutodontus*  
430 *simplex*, whose biostratigraphic range is restricted  
431 to the *Cordylodus intermedius* Zone, was originally  
432 defined in Australia (Druce & Jones, 1971) and  
433 subsequently recovered from China (Chen & Gong,  
434 1986; Chen *et al.* 1988), Laurentia (Miller, 1980; Ross  
435 *et al.* 1997; Terfelt, Bagnoli & Stouge, 2012; Miller  
436 *et al.* 2014), Siberia (Abaimova, 1971, 1975) and NW  
437 Argentina (Zeballo & Albanesi, 2009). Miller (1984)  
438 suggested that *Clavohamulus* and *Hirsutodontus* had a  
439 nektobenthic habit of life and that they preferred warm,  
440 shallow seas. The record of *Hirsutodontus* in the Los  
441 Sombreros Formation would then indicate reworking  
442 of shallow-water deposits into deep-water facies by  
443 gravity flows, as observed in the GSSP for the base of  
444 the Ordovician at Green Point, Newfoundland (Cooper,  
445 Nowlan & Williams, 2001; Miller *et al.* 2014). Ac-  
446 cordingly, *Clavohamulus hintzei* Miller, indicative of  
447 the upper subzone of *Cordylodus intermedius* Zone,  
448 is present in the shallow-marine facies of the La Silla  
449 Formation, at the eastern domain of the carbonate  
450 platform (Lehnert, Miller & Repetski, 1997).

451 The species *Teridontus nakamurai* has been found  
452 in several lithofacies suggesting a pelagic habit of life  
453 (Ji & Barnes, 1994b), eventually restricted to the shelf  
454 environments owing to a nektobenthic habit (Miller,  
455 1984). After studying a large conodont collection from  
456 the Cordillera Oriental, Zeballo & Albanesi (2013b)  
457 recognized an antithetical relationship between the cos-  
458 mopolitan genera *Variabiloconus* and *Teridontus*, veri-  
459 fying that the latter predominates in the deeper parts of  
460 the platform. In particular, *V. datsonensis* is present in  
461 NE Australia (Druce & Jones, 1971), Antarctica (Bug-  
462 gisch & Repetski, 1987) and NW Argentina (Zeballo  
463 & Albanesi, 2009).

464 The genus *Cordylodus* was a major component of  
465 most slope and platform communities during Furong-  
466 ian and Early Ordovician times. *C. proavus* has a wide-  
467 spread geographic distribution and is found in a wide  
468 range of lithofacies, which suggests a pelagic habit  
469 of life. In contrast, *C. andresi*, *C. caboti* and *C. in-*  
470 *termedius* preferred deeper-water environments (lower  
471 proximal to distal slope facies), as younger species  
472 of *Cordylodus* adapted to a nektobenthic mode of life  
473 (Miller, 1984; Zhang & Barnes, 2004).

474 The *C. intermedius* Zone has a wide global distri-  
475 bution and has been documented in China (Chen &  
476 Gong, 1986), Laurentia (Bagnoli, Barnes & Stevens,  
477 1987; Barnes, 1988; Miller, 1988; Ross *et al.* 1997;  
478 Miller *et al.* 2003) and central Asia (Dubinina, 2000).  
479 A correlative conodont assemblage has been retrieved  
480 from Australia (Druce & Jones, 1971) and Iran (Müller,  
481 1973). In Argentina, it was previously identified in  
482 the Volcancito Formation of the Famatina System (Al-  
483 banesi *et al.* 2005) and the Cardonal (Rao, 1999) and

484 Santa Rosita formations in the Cordillera Oriental (Ze-  
485 ballo & Albanesi, 2009).

486 Although conodont provinces can be already distin-  
487 guished in the late Cambrian (Jeong & Lee, 2000),  
488 most of the palaeogeographic studies are concentrated  
489 in the Ordovician, when major realms were already  
490 established by late Tremadocian times (Miller, 1984;  
491 Charpentier, 1984). Accordingly, the Precordillera is  
492 identified as a conodont faunal Province of the Temper-  
493 ate Domain of the Shallow-Sea Realm (or the Open-Sea  
494 Realm depending on the sedimentary setting) (Albanesi  
495 & Bergström, 2010; Serra & Albanesi, 2013), as it  
496 lacks the typical shallow-water, tropical forms charac-  
497 teristic of the Laurentian, Australasian or North China  
498 provinces (Bagnoli & Stouge, 1991; Zhen & Percival,  
499 2003). Instead, it is distinguished by cosmopolitan or  
500 widespread faunas, showing a moderate endemism and  
501 diversity when compared with faunas from the Tropical  
502 Domain. The Baltoscandian Province of the Cold Do-  
503 main presents lower diversities and higher abundances  
504 instead (Zhen & Percival, 2003).

505 *Macerodus diana*, an index taxon of the late  
506 Tremadocian, appeared in sample AM8A. It was first  
507 described from outcrops of the Cow Head Group in  
508 western Newfoundland, a series of Laurentian slope  
509 deposits fed from the outer shelf and the upper contin-  
510 ental slope (Fåhraeus & Nowlan, 1978; Pohler, Barnes  
511 & James, 1987). Ji & Barnes (1994a) emended its dia-  
512 gnosis with material from the Boat Harbour Forma-  
513 tion (St George Group) on the Port au Port Penin-  
514 sula in western Newfoundland. *Macerodus diana* is  
515 recognized in widely separated geographic locations  
516 of the Great Basin (Ethington & Clark, 1981; Repet-  
517 ski, 1982; Ross *et al.* 1997; Landing *et al.* 2012), the  
518 Arctic Archipelago of Canada (G. S. Nowlan, unpub.  
519 Ph.D. thesis, Univ. Waterloo, 1976) and northern Nor-  
520 way (Lehnert, Stouge & Brandl, 2013). In the Kechika  
521 Formation of British Columbia, the *Macerodus diana*  
522 Zone is absent and is substituted by the shallow-water  
523 *Scolopodus subrex* Zone (Pyle & Barnes, 2002).

524 *Rossodus* is a typical genus from the Great Basin,  
525 characteristic of the North American Midcontinent  
526 Province. The latter is approximately equivalent to  
527 the Laurentian Province of the Tropical Domain, in  
528 the Shallow-Sea Realm, distinguished by shelf areas  
529 < 200 m in depth with high endemism and diversity  
530 (Ethington & Clark, 1981; Ross *et al.* 1997; Zhen  
531 & Percival, 2003). *Rossodus manitouensis* Repetski  
532 & Ethington has also been documented in China (= *'Acodus'*  
533 *oneotensus sensu* An, Du & Gao, 1985;  
534 Wang, Bergström & Lane, 1996), Korea (Seo, Lee  
535 & Ethington, 1994), Thailand (Agematsu *et al.* 2008)  
536 and Tasmania (R. C. Cantrill, unpub. Ph.D. thesis,  
537 Univ. Tasmania, 2003). It is also known from the  
538 peri-Gondwanan volcanic arc of the Famatina Sys-  
539 tem, where it occurs along with *Drepanodus arcuatus*,  
540 *Cornuodus longibasis* (Lindström), *Paltodus deltifer*  
541 *pristinus* (Viira), *P. cf. subaequalis* Pander and *Parois-*  
542 *todus numarcuatus* (Lindström), which characterize a  
543 biofacies dominated by pelagic species from deep/cold



544 waters (Albanesi *et al.* 2005). In the eastern domain of  
 545 the Precordillera, Lehnert, Miller & Repetski (1997)  
 546 described *Rossodus* aff. *manitouensis* from shallow-  
 547 water facies of the La Silla Formation, along with *Alox-*  
 548 *oconus* cf. *propinquus* (Furnish), *Scolopodus* cf. *floweri*  
 549 Repetski, *Paroistodus numarcuatus* and *Colaptoconus*  
 550 *quadruplicatus* (Branson & Mehl). The authors correlated  
 551 this conodont assemblage with the Low Diversity  
 552 Interval and the lower *Macerodus diana*e conodont  
 553 biozone in North America (Ross *et al.* 1997), consistent  
 554 with the suggested age for the samples AM8A  
 555 and AM8B from the slope facies of the Los Sombreros  
 556 Formation.

557 The record of *Macerodus diana*e in the slope fa-  
 558 cies verifies a strong link between the conodont faunas  
 559 from the Precordillera with those from Laurentia dur-  
 560 ing Early Ordovician times, demonstrating a connec-  
 561 tion along the borders of the Iapetus Ocean. Accord-  
 562 ingly, Lehnert, Miller & Repetski (1997) interpreted  
 563 that the record of *Clavohamulus hintzei* in the La  
 564 Silla Formation as well as the faunal similarities at  
 565 the species level with the shallow-water North Amer-  
 566 ican Midcontinent Province was a consequence of the  
 567 derivation of the Cuyania Terrane from the Ouachita  
 568 Embayment in Laurentia. Nevertheless, the conodont  
 569 faunas do not provide clear evidence to postulate a  
 570 geographic origin for the Precordillera as they show  
 571 dominantly Laurentian affinities again in the Middle  
 572 Ordovician, after a gradual immigration of conodonts  
 573 from colder regions (Albanesi, 1998; Albanesi &  
 574 Bergström, 2010).

#### 575 5.d. Conodont preservation and palaeothermometry

576 The Cambrian specimens recovered from sample AM6  
 577 are well preserved and exhibit a conodont colour alter-  
 578 ation index (CAI) 3 that provides some translucency to  
 579 the conodont elements. The conodonts present smooth  
 580 surfaces and scarce mineral overgrowths. Microfrac-  
 581 tures are frequent and are responsible for the lack of  
 582 apices on cusps and denticles. Conodonts recovered  
 583 from samples AM8A and AM8B also exhibit a CAI  
 584 3 but display a sugary texture with abundant quartz  
 585 overgrowths instead. In this case, the different type of  
 586 textural alteration suggests variations in the intensity  
 587 of the diagenetic processes.

588 Interestingly, previous findings of reworked pre-  
 589 Floian conodonts recovered from the Los Sombreros  
 590 Formation display high CAI values in contrast to the  
 591 elements recovered from the host rock. This fact was  
 592 interpreted as a result of burial-related metamorphism  
 593 and exhumation of the carbonate platform near the su-  
 594 ture zone of Cuyania with Gondwana, which supplied  
 595 detritus to the deep-water basin of the Western Precor-  
 596 dillera (Voldman, Albanesi & Ramos, 2009).

597 The record of Furongian–Lower Ordovician con-  
 598 odonts with CAI 3 (~ 110–200 °C; Epstein, Epstein &  
 599 Harris, 1977) in the Los Sombreros Formation reflects  
 600 a simpler burial history instead, if a uniform palaeo-  
 601 geothermal gradient in the basin is considered. A rift-

602 related heat source is not possible to discern as CAI  
 603 values are relatively low, within the range of the ob-  
 604 served values in the platform, which can be accounted  
 605 for solely by sedimentary burial (Voldman, Albanesi &  
 606 Ramos, 2010).

607 Alternatively, the mafic rocks from the Western Pre-  
 608 cordillera produced very restricted thermal anomalies  
 609 in the country rocks given the small volume of single-  
 610 pulsed basalt intrusions, which could only slightly  
 611 contribute to a regional increment of the heat flux  
 612 (Voldman, Albanesi & do Campo, 2008; González-  
 613 Menéndez *et al.* 2013). This is consistent with the  
 614 metamorphic conditions inferred from the paragen-  
 615 etic associations of the mafic rocks, which suggests  
 616 *c.* 250–350 °C and 2–3 kbar, with palaeogeothermal  
 617 gradients of ~ 30–35 °C km<sup>-1</sup> (Robinson, Bevins &  
 618 Rubinstein, 2005), which are typical of mature pass-  
 619 ive margins and foreland basins (e.g. Allen & Allen,  
 620 2005). Consequently, the CAI 3 in the studied con-  
 621 odont samples reflects a thermal history related to the  
 622 sedimentary burial and nappe stacking of the Western  
 623 Precordillera.

#### 624 6. Conclusions

625 The new conodont data from the Los Sombreros Form-  
 626 ation in La Invernada Range along with the sedimento-  
 627 logical and structural analysis carried out in the study  
 628 area show that slope sedimentation and gravity slid-  
 629 ing has taken place in the Precordillera since at least  
 630 late Cambrian times, contrasting with the current hy-  
 631 pothesis of a Devonian age for the slope deposits of the  
 632 Los Sombreros mélange. Moreover, the recognition of  
 633 the *Hirsutodontus simplex* Subzone of the *Cordylodus*  
 634 *intermedius* Zone (upper Furongian, Cambrian) and the  
 635 *Macerodus diana*e Zone (upper Tremadocian, Ordovi-  
 636 cian) improves the correlation with the La Silla Forma-  
 637 tion of the Precordilleran carbonate platform as well as  
 638 with regions of Gondwana and other palaeocontinents.  
 639 The record of the index species *Macerodus diana*e in  
 640 the Los Sombreros Formation, as well as *Clavohamu-*  
 641 *lus hintzei* in the La Silla Formation, emphasizes the  
 642 strong faunal affinity of the shelf environments of the  
 643 Precordillera with the Laurentian Province of the Trop-  
 644 ical Domain for the late Cambrian – Early Ordovician  
 645 periods. However, given the wide global distribution of  
 646 the studied specimens, the present data are not indic-  
 647 ative of a geographic origin for the Precordillera. The  
 648 thermal alteration of the studied specimens is consist-  
 649 ent with the sedimentary burial and nappe stacking of  
 650 the Western Precordillera.

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